

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁶ : C08L 63/00, 63/02, 67/02, 71/00, 3/02, 97/02</p>	<p>A1</p>	<p>(11) International Publication Number: WO 97/23564 (43) International Publication Date: 3 July 1997 (03.07.97)</p>
<p>(21) International Application Number: PCT/US96/20194 (22) International Filing Date: 19 December 1996 (19.12.96) (30) Priority Data: 60/009,022 21 December 1995 (21.12.95) US (71) Applicant (for all designated States except US): THE DOW CHEMICAL COMPANY [US/US]; 2030 Dow Center, Mid- land, MI 48674 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): MANG, Michael, N. [US/US]; 3708 Haskin Drive, Midland, MI 48640 (US). WHITE, Jerry, E. [US/US]; 418 Huckleberry Drive, Lake Jackson, TX 77566 (US). (74) Agent: DAMOCLES, Nemia, C.; The Dow Chemical Com- pany, Patent Dept., P.O. Box 1967, Midland, MI 48641- 1967 (US).</p>		<p>(81) Designated States: BR, CA, CN, CZ, JP, KR, MX, PL, US, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p>
<p>(54) Title: COMPOSITIONS COMPRISING HYDROXY-FUNCTIONAL POLYMERS (57) Abstract Compositions comprising a polysaccharide and a hydroxy-functional polyester are prepared by mixing the hydroxy-functional polyester and polysaccharide, modified polysaccharide or a naturally-occurring fiber or particulate filler and, optionally, other additives in an intensive mixer at a temperature and for a time sufficient to provide an intimate, well-dispersed mixture of the components.</p>		

L
#5

1, 2
use for claim 1

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgyzstan	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	KZ	Kazakhstan	SG	Singapore
CH	Switzerland	LJ	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovakia
CM	Cameroon	LR	Liberia	SN	Senegal
CN	China	LT	Lithuania	SZ	Swaziland
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	LV	Latvia	TG	Togo
DE	Germany	MC	Monaco	TJ	Tajikistan
DK	Denmark	MD	Republic of Moldova	TT	Trinidad and Tobago
EE	Estonia	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	UG	Uganda
FI	Finland	MN	Mongolia	US	United States of America
FR	France	MR	Mauritania	UZ	Uzbekistan
GA	Gabon			VN	Viet Nam

COMPOSITIONS COMPRISING HYDROXY-FUNCTIONAL POLYMERS

This invention relates to compositions comprising hydroxy-functional polymers and to articles prepared therefrom.

Polysaccharides are inexpensive and have good mechanical properties, but cannot be easily processed as thermoplastics. To take advantage of their low cost, attempts have been made to blend them with synthetic polymers, such as polyolefins, to produce more useful materials. However, the resulting materials often suffer from several problems. For example, the physical properties of the blend of the polysaccharide starch with polyolefin are adversely affected by the incompatibility of and poor adhesion between the polar starch and the non-polar polyolefin. In order to solve this problem, the starch must be modified, or compatibilized with the polyolefin, thereby increasing the cost of the blend. When more polar thermoplastics are employed in the preparation of the blend, other physical properties such as moisture resistance can be adversely affected. It would be desirable to provide a composition having useful mechanical properties comprising a polysaccharide and another hydroxy-functional polymer which is compatible therewith.

The present invention is a composition comprising (1) a polysaccharide, a modified polysaccharide or a naturally-occurring fiber or particulate filler and (2) a thermoplastic hydroxy-functional polyether derived from monomers containing 1 or more epoxy groups.

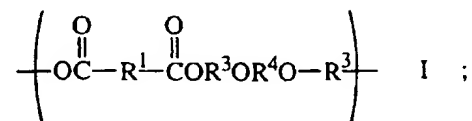
The hydroxy-functional polyether has been found to adhere strongly to a polysaccharide. This adhesion, and the generally good physical properties of hydroxy-functional polyethers, allows for the preparation of hitherto unknown materials with useful physical properties.

The compositions of the present invention are suitable for use in fabricating molded, extruded or foamed articles, containers, films, film laminates, or coatings using conventional fabricating techniques such as extrusion, compression molding, injection molding, blow molding and similar fabrication techniques commonly used to produce such articles. Examples of such articles include films, foams, sheets, pipes, rods, bags, boxes, meat trays, egg cartons, cups and plates, cutlery,

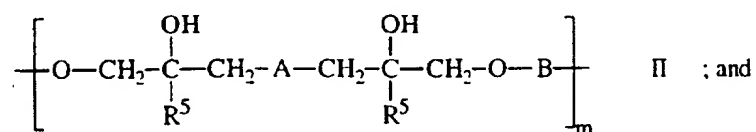
and other disposable consumer items. The compositions of the present invention are also suitable for use as adhesives and encapsulating agents.

Preferably, the hydroxy-functional polyethers employed in the practice of the present invention are:

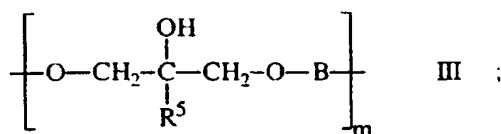
- 5 (1) hydroxy functional polyethers having repeating units represented by the formula:



- (2) hydroxy-functional polyethers having repeating units represented by the formula:

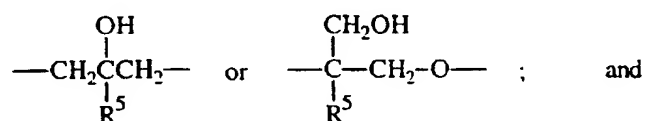


- (3) hydroxy-functional polyethers having repeating units represented by the formula:

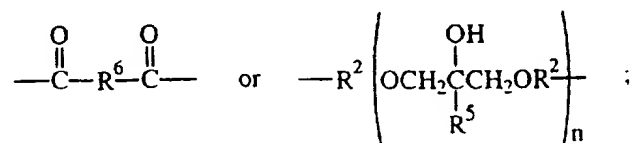


10

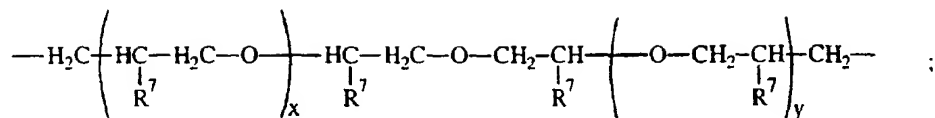
wherein R¹ individually represents a divalent organic moiety which is predominantly hydrocarbylene, or a combination of different organic moieties which are predominantly hydrocarbylene; R³ is



- 15 R⁴ is



wherein R² is a divalent organic moiety which is predominantly hydrocarbylene or



- R^5 is hydrogen or alkyl, R^6 is independently an organic moiety which is predominantly hydrocarbylene, R^7 is independently hydrogen or methyl, A is an amine moiety or a combination of different amine moieties; B is a divalent organic moiety which is predominantly hydrocarbylene; m is an integer from 10 to about 1000, n is an integer from about 0 to about 100, and x and y are independently integers from 0 to 100.

- The term "predominantly hydrocarbylene" is defined as a divalent radical that is predominantly hydrocarbon, but which optionally contains a minor amount of a heteroatomic moiety such as oxygen, sulfur, imino, sulfonyl, and sulfoxyl.

- Representative divalent organic moieties useful as R^1 , R^2 and R^6 include alkylene, cycloalkylene, alkylenearylene, poly(alkyleneoxyalkylene), alkyleneethioalkylene, alkylene sulfonylalkylene, alkylene substituted with at least one hydroxyl group, cycloalkylene substituted with at least one hydroxyl group, alkylenearylene substituted with at least one hydroxyl group, poly(alkyleneoxyalkylene) substituted with at least one hydroxyl group, alkyleneethioalkylene substituted with at least one hydroxyl group, alkylene sulfonylalkylene substituted with at least one hydroxyl group; arylene, dialkylenearylene, diaryleneketone, diarylenesulfone, diarylene oxide and diarylene sulfide.

- In the more preferred hydroxy-functional polyethers, R^1 , R^2 and R^6 are independently methylene, ethylene, propylene, butylene, pentamethylene, hexamethylene, heptamethylene, octamethylene, nonamethylene, decamethylene, dodecamethylene, 1,4-cyclohexylene, 1,3-cyclohexylene or 1,2-cyclohexylene optionally substituted with at least one hydroxyl group, p-phenylene, m-phenylene, or 2,6-naphthalene, diphenylene-isopropylidene, sulfonyldiphenylene, carbonyldiphenylene, oxydiphenylene, or 9,9-fluorenediphenylene and n is from 0 to 10.

Preferably, A is 2-hydroxyethylimino-, 2-hydroxypropylimino-, piperazanyl, N,N'-bis(2-hydroxyethyl)-1,2-ethylenediimino-, and B is isopropylidenediphenylene, 1,3-phenylene, or 1,4-phenylene, and R⁵ is hydrogen.

The poly(hydroxy ester ethers) represented by Formula I are prepared
5 by reacting diglycidyl esters of aliphatic or aromatic diacids such as diglycidyl terephthalate, or diglycidyl ethers of dihydric phenols or alcohols with aliphatic or aromatic diacids such as adipic or terephthalic acid as described in U.S. Patents 5,171,820 and 5,496,910; and copending U.S. Patent Application Serial Nos. 131,110, filed October 1, 1993; 278,361, filed July 21, 1994; and 280,425, filed
10 July 26, 1994.

The poly(hydroxy amino ethers) represented by Formula II are prepared by contacting one or more of the diglycidyl ethers of a dihydric phenol with an amine having two amine hydrogens under conditions sufficient to cause the amine moieties to react with epoxy moieties to form a polymer backbone having amine
15 linkages, ether linkages and pendant hydroxyl moieties. These polyethers are described in U.S. Patent 5,275,853. These polyethers can also be prepared by contacting a diglycidyl ether or an epihalohydrin with a difunctional amine.

The poly(hydroxy ethers) represented by Formula III can be prepared by contacting a diglycidyl ether or a combination of diglycidyl ethers with a dihydric
20 phenol or combination of dihydric phenols using the process described in U. S. Patent 5,164,472. Alternatively, the poly(hydroxy ethers) are obtained by allowing a dihydric phenol or a combination of dihydric phenols to react with an epihalohydrin by the process described by Reinking, Barnabeo, and Hale in the Journal of Applied Polymer Science, Volume 7, page 2135 (1963).

25 While the amount of hydroxy-functional polyethers used depends on a variety of factors, including the specific polymer employed and the desired end uses of the composition, in general, the hydroxy-functional polyesters can be present in an amount of from 1 to 99 weight percent, preferably from 10 to 95 weight percent and, most preferably from 20 to 90 weight percent, based on the total weight of the
30 hydroxy-functional polyester and polysaccharide.

The naturally-occurring fibers or particulate fillers which can be employed in the practice of the present invention for preparing the composition are,

for example, wood flour, wood pulp, wood fibers, cotton, flax, hemp, or ramie fibers, rice or wheat straw, chitin, chitosan, cellulose materials derived from agricultural products, nut shell flour, corn cob flour, and mixtures thereof. Polysaccharides which can be employed in the practice of the present invention for preparing the

5 composition are the different starches, celluloses, hemicelluloses, xylanes, gums, pectins and pullulans. Polysaccharides are known and are described, for example, in Encyclopedia of Polymer Science and Technology, 2nd edition, 1987. The preferred polysaccharides are starch and cellulose.

Modified polysaccharides which can be employed in the practice of the
10 present invention for preparing the composition are the esters and ethers of polysaccharides, such as, for example, cellulose ethers and cellulose esters, or starch esters and starch ethers. Modified polysaccharides are known and are described, for example, in Encyclopedia of Polymer Science and Technology, 2nd edition, 1987

15 The term "starch" as used herein, refers to carbohydrates of natural, vegetable origin, composed mainly of amylose and/or amylopectin, and includes unmodified starches, physically modified starches, such as thermoplastic, gelatinized or cooked starches, starches with a modified acid value (pH) where acid has been added to lower the acid value of a starch to a range of from 3 to 6, gelatinized
20 starches, ungelatinized starches, cross-linked starches and disrupted starches (starches which are not in particulate form). The starches can be in granular, particulate or powder form. They can be extracted from various plants, such as, for example, potatoes, rice, tapioca, corn, pea, and cereals such as rye, oats, and wheat.

25 Celluloses are known and are described, for example, in Encyclopedia of Polymer Science and Technology, 2nd edition, 1987. Celluloses are natural carbohydrate high polymers (polysaccharides) consisting of anhydroglucose units joined by an oxygen linkage to form long molecular chains that are essentially linear. Cellulose can be hydrolyzed to form glucose. The degree of polymerization
30 ranges from 1000 for wood pulp to 3500 for cotton fiber, giving a molecular weight of from 160,000 to 560,000. Cellulose can be extracted from vegetable tissues (wood, grass, and cotton). Celluloses can be used in the form of fibers.

The composition of the present invention may also contain various additives such as, for example, plasticizers, lubricants, pigments, foaming agents, extenders, stabilizers, chemical modifiers, and flow accelerators. Each of these additives is known and several types of each are commercially available.

5 In general, the compositions of the present invention can be prepared by mixing (1) a thermoplastic hydroxy-functional polyether and (2) a polysaccharide, a modified polysaccharide or a naturally-occurring fiber or particulate filler and, optionally (3) other additives in an intensive mixer, such as a Haake mixer, a Banbury mixer, single- screw extruder, twin-screw extruder, or an injection-molding machine,
10 at a temperature and for a time sufficient to provide an intimate, well-dispersed mixture of the components. Preferably, the components are brought together and processed in an appropriate melt extruder from which the blend is extruded in the form of powder or strands. The powders or strands are then pelletized or formed into granules for injection molding and other thermal processes. Standard techniques
15 and apparatuses well known in the art can be used for these processes.

 The material can be fabricated into films using processes well known to those skilled in the art, such as by cast film extrusion, blown film extrusion, and can be incorporated in multilayer constructions such as coextruded cast or blown films, with or without intermediate adhesive layers or layers containing recycled scrap
20 material from the fabrication process.

 The material can be extruded into shapes such as profiles, tubes, rods, strips, tapes, sheets, or strands, and may be further formed by molding using well known methods such as compression or injection molding, or thermoforming or vacuum molding.

25 The material may be in the form of a foam or cellular structure, as a flexible, rigid, or structural foam, with open, closed, or partially opened-cellular morphologies. The foam can be conveniently formed by extrusion, or by expansion of beads into a heated mold to form a shaped article. The extruded foam may be further formed by thermoforming into shaped articles. The foam may also be directly
30 extruded into useful shapes such as rods, slabs, planks, or boards. The articles and shapes so formed are useful, for example, for packaging, insulation, and cushioning purposes.

The foam of the present invention is generally prepared by heating the composition comprising a polysaccharide, a modified polysaccharide, a naturally occurring fiber or particulate filler, and a hydroxy-functional polyether to form a plasticized or molten polymer material, incorporating a blowing agent to form a foamable gel, and extruding the gel through a die to form the foam product. The blowing agent can be incorporated into the polymer material before melting, or may be incorporated after the polymer material is heated to a temperature at or above its melting point. The blowing agent may be incorporated or mixed into the melt polymer material by any means known in the art such as with an extruder, mixer, or blender.

10 The blowing agent is mixed with the melt polymer material at an elevated pressure sufficient to prevent substantial expansion of the melt polymer material and to generally disperse the blowing agent homogeneously therein. Optionally, a nucleator may be blended in the polymer melt or dry blended with the polymer material prior to plasticizing or melting. The foamable gel is typically cooled to a lower temperature to

15 optimize physical characteristics of the foam structure. The gel is then extruded through a die of desired shape to a zone of lower or reduced pressure than that in which the gel is maintained prior to extrusion through the die. The lower pressure may be superatmospheric or subatmospheric, but is preferably at an atmospheric level.

20 Blowing agents useful in making the present foam structures include inorganic agents, organic blowing agents, and chemical blowing agents. Suitable inorganic blowing agents include carbon dioxide, nitrogen, argon, water, air, and helium. Organic blowing agents include aliphatic hydrocarbons having 1 to 9 carbon atoms and fully and partially halogenated aliphatic hydrocarbons having 1 to 4 carbon

25 atoms.

The present foam has the density from 5 to 200 kilograms per cubic meter. The foam has an average cell size from 0.1 to 5.0 millimeters. The foam may be open or closed-cell.

Though the preferred process for making the present foam is an extrusion process, it is understood that the above structure may be formed by the expansion of beads, which may be molded at the time of expansion to form structures of various shapes.

30

The composition of the present invention is also useful as hot-melt adhesives, and can be formulated with tackifying resins, plasticizers, waxes, and/or conventional additives in varying amounts as are known to those skilled in the art.

5 The composition may be fabricated into a container using conventional processes such as blow molding, injection molding, vacuum forming, thermoforming, injection blow molding, extrusion blow molding, and pultrusion.

The composition can be used as a coating or a laminate, and can be applied to a substrate by methods such as calendering, curtain coating, extrusion coating, roll coating, or spray coating.

10 The composition can be used as an encapsulant capable of slow or controlled release of a pharmacologically active agent, a catalyst, a biocide or a fertilizer, and can be prepared by compounding the material with, as an additional component, the pharmacologically active agent, the biocide, the fertilizer, or the fertilizer, either during or after the composition is prepared. In the case of active
15 materials that might not be stable under the processing conditions employed to produce the composition, the composition of the invention can be applied to particles of the active agent by spray coating, solution coating, or other well known methods to produce the encapsulated active ingredient.

20 The composition may also be in the form of a substrate comprising a polysaccharide, such as wood, paper, textile, or a starch article, optionally containing a hydroxy-functional polyether derived from monomers containing one or more epoxy groups, coated with a hydroxy-functional polyether derived from monomers containing one or more epoxy groups. The hydroxy-functional polyether can be used to modify the surface or structural properties of the substrate, to protect the substrate
25 in use. The coating can be conveniently applied using well known methods such as extrusion, calendering, pressure lamination, roll coating, powder coating, curtain coating, or solution coating. The substrate is preferably a paper, cardboard, cellulose film, modified cellulose film, starch film, a modified starch film, wood, or a film or an article comprising a polysaccharide, a modified polysaccharide, a naturally occurring
30 fiber or particulate filler, and a hydroxy-functional polyether. The substrate is most preferably paper, cardboard, or an article comprising a polysaccharide, a modified

polysaccharide, a naturally occurring fiber or particulate filler, and a hydroxy-functional polyether.

The following working examples are given to illustrate the invention and should not be construed as limiting its scope. Unless otherwise indicated, all parts and percentages are by weight.

Example 1

A 13 L resin kettle, equipped with a mechanical stirrer and nitrogen inlet, was charged with hydroquinone diglycidyl ether (1312.5 g, 5.8 mol, 113.06 g/equivalent epoxide), 1,10-decanedicarboxylic acid (1343.5 g, 5.83 mol), and tetra-n-butylammonium bromide (94.2 g, 0.29 mol). Diglyme (3 L) was added and the mixture heated to 110°C for 5.5 hours under an atmosphere of nitrogen. Glacial acetic acid (250 mL) was added and heating at 110°C was continued overnight. The solution was allowed to cool to about 50°C and was poured into water in a large Waring blender (300 mL portions into 2 L of water). The fibrous precipitate was collected by suction filtration and suspended in fresh water for 3 days. The product was collected by suction filtration and allowed to air-dry overnight. The product is dried in a vacuum oven at 100°C to 115°C overnight. The polymer had an inherent viscosity of 0.42 dL/g (DMF, 25°C, 0.5g/dL), a T_g of 5°C, and a T_m of 75°C.

Dry blends of the above polymer and unmodified potato starch were prepared using the weight ratios indicated in the table. The blends were compounded using a Haake mixer (60 cc bowl) at 120°C for 6 minutes as indicated. Plaques (4 inch by 4 inch by 1/16 inch) were prepared by compression molding the material obtained from the Haake mixer. Specimens for mechanical property testing were obtained from these plaques. Selected tensile properties are listed in Table I.

Table IMechanical Properties of Starch-Poly(ester ether) Blends

<u>Mech. Properties</u>	<u>Control (0% Starch)</u>	<u>20% Starch</u>	<u>40% Starch</u>	<u>60% Starch</u>	<u>80% Starch</u>
Yield Stress (PSI)	880	940	1130	1340	-
Yield Strain (%)	15	12	9	8	-
Tensile Strength (psi)	3050	1840	870	1080	2210
Tensile Elongation (%)	705	370	190	14	6
Tensile Modulus (psi)	12240	16640	29690	45210	96510

Example 2

- 5 The poly(hydroxy amino ether) derived from the reaction of the diglycidyl ether of bisphenol A with ethanolamine was compounded with varying amounts of hardwood flour (American Wood Fibers grade 20010) using a mechanical Brabender Plasticorder with roller blades in a nominal 60 cc counter-rotating mixer head at 180°C. The polymer was loaded and processed at 63 rpm for two minutes,
- 10 then the hardwood flour was added and processed under the conditions shown in Table II.

Table II

<u>Example number</u>	<u>Weight % wood flour</u>	<u>Minutes processed</u>	<u>Final torque (m-g)</u>
2(a)	10	5	2500
2(b)	30	3	3200
2(c)	50	2	4500

The compounded materials were compression-molded into test plaques using a 4 inch by 4 inch by 1/16 inch thick frame mold chase between two flat stainless steel plates using Fluoroglide™ CP as a mold release. The material was molded at 200°C for 3 minutes using 1250 psi on the mold cavity. The plaque was cooled under pressure to below 45°C before demolding. ASTM Type IV tensile test bars were machined from these plaques and tested using an Instron 4507 testing frame at a crosshead speed of 0.2 inch/minute. The test results are shown in Table III.

Table III

<u>Example number</u>	<u>Tensile strength at break (psi)</u>	<u>% Elongation at break</u>	<u>Tensile modulus (psi)</u>
2(a)	6500	10.6	522,000
2(b)	10,300	3.5	670,000
2(c)	11,100	2.2	813,000

10

Example 3

The poly(hydroxy ester ether) derived from the reaction of the diglycidyl ether of bisphenol A with adipic acid was compounded with various levels of hardwood flour (American Wood Fibers grade 20010) using a mechanical Brabender Plasticorder with roller blades in a nominal 60 cc counter-rotating mixer head at 150°C. The polymer was loaded and processed at 63 rpm for two minutes, then the wood flour was added and processed under the conditions shown in Table IV.

Table IV

<u>Example number</u>	<u>Weight % wood flour</u>	<u>Minutes processed</u>	<u>Final torque (m-g)</u>
3(a)	10	3	400
3(b)	30	5	520
3(c)	50	5	1500

The compounded materials were compression-molded into test plaques using a 4 inch by 4 inch by 1/16 inch thick frame mold chase between two flat stainless steel plates using Fluoroglide™ CP as a mold release. The Examples 3(a) and 3(b) materials were molded at 150°C for 3 minutes using 1125-
 5 1250 psi on the mold cavity. The Example 3 (c) material was molded at 175°C. The plaques were cooled under pressure to below 30°C before demolding. ASTM Type IV tensile test bars are machined from these plaques and tested using an Instron 4507 testing frame at a crosshead speed of 0.2 inch/minute. The results of the test are shown in Table V.

10

Table V

<u>Example number</u>	<u>Tensile strength at break (psi)</u>	<u>% Elongation at break</u>	<u>Tensile modulus (psi)</u>
3(a)	3300	13.6	451,000
3(b)	5700	4.5	648,000
3(c)	7800	2.0	843,000

Example 4

The poly(hydroxy ether) of bisphenol A (35 g, PAPHEN™ PHENOXY RESIN PKHHMM from Phenoxy Associates) was compounded with 15 grams of
 15 dried corn starch using a Haake Rheocord 9000 torque rheometer equipped with a Haake Model 600 mixer using roller blades at a bowl temperature of 160°C. The polymer and starch were loaded and processed at 60 rpm for ten minutes. The torque and melt temperature equilibrated at a torque of about 1650 m-g at 183°C after about 5 minutes.

20

The compounded material was compression-molded into test plaques using a 4 inch by 4 inch by 1/16 inch thick frame mold chase between two flat stainless steel plates using Fluoroglide™ CP (a product of Norton Performance Plastic Company) an aerosol fluorocarbon mold release agent. The material was
 25 molded at 200°C for 3 minutes using 625 psi on the mold cavity. The plaque was then cooled under pressure to below 45°C before demolding. ASTM Type IV tensile

test bars were machined from these plaques and tested as in Example 1. The test results are shown in Table VI

Table VI

<u>Tensile strength at break</u> <u>(psi)</u>	<u>% Elongation at break</u>	<u>Tensile modulus</u> <u>(psi)</u>
8431	2.47	499,600

5 Example 5

The poly(hydroxy amino ether) derived from the reaction of the diglycidyl ether of bisphenol A with ethanolamine (35 g) was compounded with 15 grams of dried corn starch (Buffalo 3401) using a Haake Rheocord 9000 torque rheometer equipped with a Haake Model 600 mixer using roller blades at a bowl
10 temperature of 160°C. The polymer and starch were loaded and processed at 60 rpm for ten minutes. The torque and melt temperature equilibrated at a torque of about 1500 m-g at 183°C after about 5 minutes.

The compounded material was compression -olded into test plaques using a 4 inch by 4 inch by 1/16 inch thick frame mold chase between two flat
15 stainless steel plates using Fluoroglide™ CP as a mold release agent. The material was molded at 200°C for 3 minutes using 625 psi on the mold cavity. The plaque was then cooled under pressure to below 45°C before demolding. ASTM Type IV tensile test bars are machined from these plaques and tested as in Example 1. The test results are shown in Table VII.

20

Table VII

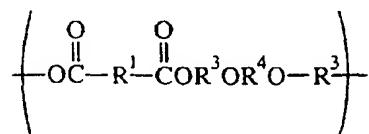
<u>Tensile strength at break</u> <u>(psi)</u>	<u>% Elongation at break</u>	<u>Tensile modulus</u> <u>(psi)</u>
8134	2.75	551,000

The data above indicate that addition of starch to a hydroxy-functional polyether in accordance with the present invention produces materials having useful mechanical properties.

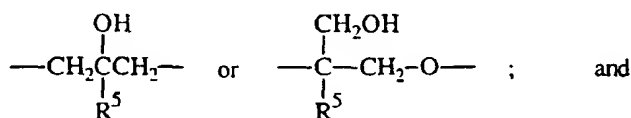
CLAIMS:

1. A composition of matter comprising (1) a polysaccharide, a modified polysaccharide, or a naturally-occurring fiber or particulate filler and (2) a thermoplastic hydroxy-functional polyether derived from monomers containing 1 or more epoxy groups.

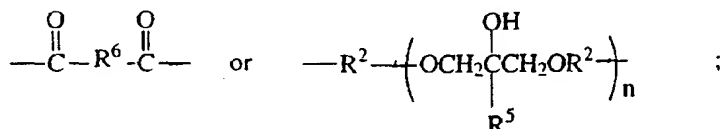
2. The composition of Claim 1, wherein the thermoplastic hydroxy-functional polyether has repeating units represented by the formula:



wherein R¹ is a divalent organic moiety which is predominantly hydrocarbylene; R³ is



R⁴ is



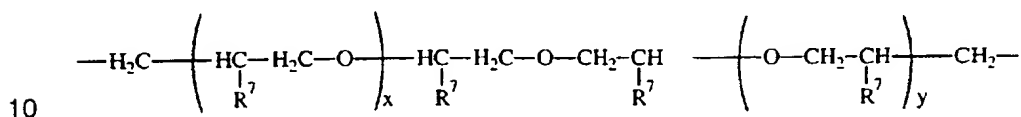
wherein R² and R⁶ are independently divalent organic moieties which are predominantly hydrocarbon, R⁵ is hydrogen or alkyl; and n is from 0 to 100.

3. The composition of Claim 2, wherein R¹, R² and R⁶ are independently alkylene, cycloalkylene, alkylenearylene, poly(alkyleneoxyalkylene), alkyleneethioalkylene, alkylene-sulfonylalkylene, alkylene substituted with at least one hydroxyl group, cycloalkylene substituted with at least one hydroxyl group, alkylenearylene substituted with at least one hydroxyl group, poly(alkyleneoxyalkylene) substituted with at least one hydroxyl group, alkyleneethioalkylene substituted with at least one hydroxyl group, alkylene-sulfonylalkylene substituted with at least one hydroxyl group, arylene,

dialkylenearylene, diaryleneketone, diarylenesulfone, diarylene oxide or diarylene sulfide.

4. The composition of Claim 2, wherein R¹, R² and R⁶ are independently methylene, ethylene, propylene, butylene, pentamethylene, hexamethylene, heptamethylene, octamethylene, nonamethylene, decamethylene, dodecamethylene, 1,4-cyclohexylene, 1,3-cyclohexylene or 1,2-cyclohexylene optionally substituted with at least one hydroxyl group.

5. The composition of Claim 2, wherein R¹, R² and R⁶ are represented by the formula:



wherein R⁷ is independently hydrogen or methyl and x and y are independently from 0 to 100.

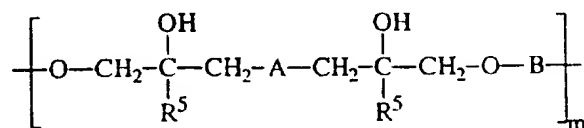
6. The composition of Claim 2, wherein R¹ and R⁶ are independently p-phenylene, m-phenylene, or 2,6-naphthalene.

7. The composition of Claim 2, wherein R² is m-phenylene, p-phenylene, naphthylene, diphenylene-isopropylidene, sulfonyldiphenylene, carbonyldiphenylene, oxydiphenylene, or 9,9-fluorenediphenylene.

8. The composition of Claim 2, wherein R⁵ is hydrogen.

9. The composition of Claim 2, wherein n is from 0 to 10.

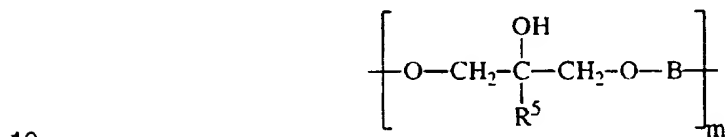
10. The composition of Claim 1, wherein the thermoplastic hydroxy-functional polyether has repeating units represented by the formula:



wherein A is an amine moiety or a combination of different amine moieties; B is a divalent organic moiety which is predominantly hydrocarbylene; R⁵ is alkyl or hydrogen; and m is an integer from 10 to 1000.

11. The composition of Claim 10, wherein A is 2-hydroxyethylimino-, 2-hydroxypropylimino-, piperazeryl, N,N'-bis(2-hydroxyethyl)-1,2-ethylenediimino-, and B is isopropylidenediphenylene, 1,3-phenylene, or 1,4-phenylene, and R⁵ is hydrogen.

12. The composition of Claim 1, wherein the thermoplastic hydroxy-functional polyether is represented by the formula:



wherein B is a divalent organic moiety which is predominantly hydrocarbylene, and R⁵ is alkyl or hydrogen, and m is an integer from 10 to 1000.

13. The composition of Claim 12, wherein B is isopropylidenediphenylene, 1,3-phenylene, or 1,4-phenylene, and R⁵ is hydrogen.

14. The composition of Claim 12, wherein the thermoplastic hydroxy-functional polyether is derived from the reaction of an epihalohydrin and a bisphenol.

15. The composition of Claim 1, wherein the polysaccharide is a starch or a cellulose.

16. The composition of Claim 15, wherein the starch is derived from potato, rice, corn or wheat.

17. The composition of Claim 15, wherein the starch is a granular starch, and the cellulose is in the form of a fiber.

18. The composition of Claim 15, wherein the starch is a thermoplastic starch.

19. The composition of Claim 15, wherein the cellulose is extracted from wood pulp or cotton fibers.

20. The composition of Claim 1, wherein the modified polysaccharide is an ether or an ester of the polysaccharide.

5 21. The composition of Claim 20, wherein the modified polysaccharide is a cellulose ether or cellulose ester.

22. The composition of Claim 1, wherein the naturally-occurring fiber or particulate filler is wood flour, wood pulp, wood fibers, cotton, flax, hemp, or ramie fibers, rice or wheat straw, chitin, chitosan, cellulose materials derived from
10 agricultural products, nut shell flour, corn cob flour, or mixtures thereof.

23. The composition of Claim 1, wherein the thermoplastic hydroxy-functional polyether is present in an amount of from 1 to 99 weight percent, and the polysaccharide, a modified polysaccharide, or a naturally-occurring fiber or particulate filler is present in an amount of from 99 to 1 weight percent, based on the weight of
15 the two components.

24. The composition of Claim 1, in the form of a film.

25. The composition of Claim 1, in the form of a molded or extruded article.

26. The composition of Claim 1, in the form of an open-cell or closed
20 cell foam.

27. The composition of Claim 1, in the form of an adhesive.

28. The composition of Claim 1, in the form of a rigid or flexible container.

29. The composition of Claim 1, in the form of a packaging material.

25 30. The composition of Claim 1, in the form of a coating.

31. The composition of Claim 1, in the form of a laminate.

32. The composition of Claim 1, in the form of an encapsulant capable of slow or controlled release of a pharmacologically active agent, a catalyst, a biocide or a fertilizer.

33. An article comprising a substrate of a polysaccharide coated with a thermoplastic hydroxy-functional polyether derived from monomers containing 1 or more epoxy groups.

34. The article of Claim 33, in which the polysaccharide is a paper product.

35. A laminate comprising a thermoplastic hydroxy-functional polyether derived from monomers containing 1 or more epoxy groups bonded to a substrate comprising, paper, cardboard, cellulose film, modified cellulose film, starch film, modified starch film, or wood.

36. A process for preparing the composition of Claim 1, which comprises mixing the thermoplastic hydroxy-functional polyether derived from monomers containing 1 or more epoxy groups with the polysaccharide, modified polysaccharide, or naturally-occurring fiber or particulate filler in an intensive mixer at a temperature and for a time sufficient to provide a well-dispersed mixture of the components.

INTERNATIONAL SEARCH REPORT

In tional Application No
PCT/US 96/20194

A. CLASSIFICATION OF SUBJECT MATTER IPC 6 C08L63/00 C08L63/02 C08L67/02 C08L71/00 C08L3/02 C08L97/02		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 6 C08L		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4 179 540 A (SMAROOK W.A.) 18 December 1979 see column 8, line 24 - line 27; table 1 see column 8, line 38 - line 45 see column 3, line 32 - column 4, line 10 see claims 1,8,26,29 ---	1,12-15, 17,19, 23,25, 26,28,31
X	DATABASE WPI Section Ch, Week 7625 Derwent Publications Ltd., London, GB; Class A85, AN 76-46762X XP002030040 & JP 51 050 708 A (SONY CORP) , 4 May 1976 see abstract --- -/--	1,12-15, 23,24
<input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "Z" document member of the same patent family		
Date of the actual completion of the international search 24 April 1997		Date of mailing of the international search report 26.05.97
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer O'Sullivan, T

INTERNATIONAL SEARCH REPORT

Int. Patent Application No.
PCT/US 96/20194

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>DATABASE WPI Section Ch, Week 8344 Derwent Publications Ltd., London, GB; Class A25, AN 83-804399 XP002030041 & JP 58 160 368 A (SUMITOMO CHEM CO LTD) , 22 September 1983 see abstract ---</p>	<p>1,12,14, 15,20, 23,27,30</p>
X	<p>GB 1 014 884 A (HAWLEY PRODUCTS) 30 July 1962 see examples 1,16,17,23 ---</p>	<p>1,10, 15-17, 22,23, 33-35</p>
X	<p>US 3 313 640 A (BAUER K.) 11 April 1967 see column 1, line 15 - line 20 see example 1 ---</p>	<p>1,15-17, 23,33,36</p>
A	<p>EP 0 513 679 A (DOW CHEMICAL CO) 19 November 1992 cited in the application see claims 1-16 -----</p>	<p>1-10</p>

INTERNATIONAL SEARCH REPORT

In .tional Application No

PCT/US 96/20194

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4179540 A	18-12-79	US 4127547 A	28-11-78
		CA 1080897 A	01-07-80
		DE 2558044 A	08-07-76
		FR 2295828 A	23-07-76
		GB 1505061 A	22-03-78
		GB 1504559 A	22-03-78
		JP 1210595 C	12-06-84
		JP 51089566 A	05-08-76
		JP 58036004 B	06-08-83

GB 1014884 A		BE 621064 A	
		DE 1520057 A	26-03-70
		FR 1348382 A	06-04-64
		US 3336247 A	15-08-67

US 3313640 A	11-04-67	NONE	

EP 0513679 A	19-11-92	US 5171820 A	15-12-92
		AU 643173 B	04-11-93
		AU 1618192 A	19-11-92
		CA 2068497 A	14-11-92
		JP 5306332 A	19-11-93
